

LEMBAR
HASIL PENILAIAN SEJAWAT SEBIDANG ATAU PEER REVIEW
KARYA ILMIAH: JURNAL ILMIAH

Judul Karya Ilmiah (artikel) : Preparation, Characterization, and Activation of Co-Mo/Y Zeolite Catalyst for Coal Tar Conversion to Liquid Fuel

Jumlah Penulis : 4 orang

Status Pengusul : Penulis Pertama

Identitas Buku : a. Nama Jurnal : Bulletin of Chemical Reaction Engineering and Catalysis
b. Nomor ISSN : 1978-2993
c. Volume, nomor, bulan, tahun : Vol. 12, No. 2, p.219-226 (Februari 2017)
d. Penerbit : Teknik Kimia, Universitas Diponegoro
e. DOI artikel (jika ada) : <https://doi.org/10.9767/bcrec.12.2.768.219-226>
f. Alamat URL Jurnal/Artikel : <https://ejournal2.undip.ac.id/index.php/bcrec/article/view/768>
g. Terindeks di (Impact Factor) : SCOPUS, DOAJ, EBSCO (SJR = 0.251)

Kategori Publikasi Karya Ilmiah : ☒ Jurnal Ilmiah Internasional/ Internasional Bereputasi.

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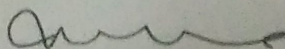
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Semarang, 16 Agustus 2018

Reviewer 2



Prof. Dr. Abdullah, MS

NIP. 19551231 198303 1 014

(Bidang Ilmu/Unit kerja : Teknik Kimia/Universitas Diponegoro)

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Komponen Yang Dinilai	Nilai Maksimum Jurnal			Nilai Akhir Yang Diperoleh
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a. Kelengkapan unsur isi jurnal (10%)	4			4,0
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- Data yang dipakai cukup lengkap dan mutakhir serta metodologi yang digunakan baik
- Kualitas penerbit cukup baik oleh Teknik Kimia Universitas Diponegoro, ISSN 1978-2993, Indexed by: SCOPUS, DOAJ, EBSCO (Q-4)
- Keterbaruan artikel ini adalah menggunakan katalis Co-Mo/zeolit Y untuk konversi coal tar, yang merupakan limbah industri

Semarang, 16 Agustus 2018

Reviewer 1

Prof. Dr. Ir. Bakti Jos, DEA

NIP. 196005011986031003

(Bidang Ilmu/Unit kerja : Teknik Kimia/Universitas Diponegoro)

PREPARATION, CHARACTERIZATION and ACTIVATION of Co-Mo/Y ZEOLITE CATALYST for CONVERSION of COAL TAR to LIQUID FUEL

Didi Dwi Anggoro*, Luqman Buchori, Giveni Christina S., Resti Nur U.

Department of Chemical Engineering, University of Diponegoro
Jl. Prof. Soedarto, Tembalang, Semarang, Indonesia 50275

* Corresponding author, tel/fax: 6224-76480675, email: anggorophd@gmail.com
Co-author 1, email: luqman.buchori@gmail.com
Co-author 2, email: givenics@gmail.com
Co-author 3, email: nurutamiresti48@gmail.com

ABSTRACT

One of many efforts to convert coal tar into alternative liquid fuel is by hydrocracking. This research aims to determine the impregnation of Co-Mo/Y zeolite, its characteristics, the effect of impregnation temperature and time, and also the best Co-Mo/Y zeolite impregnation condition for the conversion of coal tar. This research was conducted in several steps, impregnating Co in $\text{Co}(\text{NO}_3)_2 \cdot 6\text{H}_2\text{O}$ and Mo in $(\text{NH}_4)_6\text{Mo}_7\text{O}_{24} \cdot 4\text{H}_2\text{O}$ into Y zeolite in liquid media, drying at 100°C for 24 hours, and calcination at 550°C for 3 hours. Coal tar was then reacted with Co-Mo/Y zeolite and H_2 flowing was executed at 350°C . Characteristic analysis showed that Co and Mo had impregnated into Y zeolite, as well as it made no change of catalyst's structure and increased the total acidity. The higher the impregnation temperature was, the higher catalyst crystallinity, total acidity, and yield were. The longer impregnation took time, the more crystallinity reduced but total acidity and yield increased. GC analysis showed that product in the time range of gasoline contained C_8 , C_9 , and C_{10} .

Keywords: *impregnation; Co-Mo/Y zeolite; characterization; coal tar; liquid fuel*

ABSTRAK

Satu usaha untuk mengkonversi tar batubara menjadi bahan bakar cair alternative dengan proses perengkahan-hidro. Tujuan dari penelitian ini adalah membuat katalis Co-Mo/Y zeolit secara impregnasi, melakukan karakterisasi, mengetahui pengaruh suhu dan waktu impregnasi, serta menentukan kondisi optimum untuk mengkonversi tar batubara. Penelitian ini dijalankan dengan beberapa langkah, yaitu : mengimpregnasi Co dari $\text{Co}(\text{NO}_3)_2 \cdot 6\text{H}_2\text{O}$ dan Mo dari $(\text{NH}_4)_6\text{Mo}_7\text{O}_{24} \cdot 4\text{H}_2\text{O}$ kedalam zeolit Y fase cair, pengeringan pada suhu 100°C selama 24 jam, dan kalsinasi pada suhu 550°C selama 3 jam. Tar batubara kemudian direaksikan dengan katalis Co-Mo/Y zeolit dengan dialiri gas H_2 pada suhu 350°C . Hasil dari karakterisasi menunjukkan bahwa impregnasi Co dan Mo kedalam zeolit Y tidak mengubah struktur katalis tetapi merubah persen kristallinitas dan derajat keasaman. Semakin besar suhu imregnasi menyebabkan persen kristallinitas, derajat keasaman dan yield produk meningkat. Semakin lama waktu impregnasi menyebabkan persen kristallinitas menurun, tetapi derajat keasaman dan yield produk meningkat. Analisa GC memperlihatkan bahwa produk cair yang dihasilkan termasuk bensin range, yang mengandung C_8 , C_9 , dan C_{10} .

Kata Kunci: *impregnasi; Co-Mo/Y zeolit; karakterisasi; tar batubara; bahan bakar cair*

INTRODUCTION

The use of petroleum as a fuel in Indonesia is increasing every year. While oil production constantly decreased. To overcome this, the government officially issued Regulation No. 5 Year 2006 on National Energy Policy where in it is mentioned to reduce petroleum consumption and increase the consumption of other energies such as coal.

Coal should not be directly burned, but would be more meaningful and efficient if converted into synthetic fuel or other high value petrochemical materials. One is by a gasification process. Coal gasification is a process to convert solid coal into flammable coal gas. This process produces byproducts such as coal tar which is quite large. Coal tar is black or dark brown high viscosity liquid. Coal tar has very complex chemical components, includes monocyclic aromatic compounds, polycyclic aromatic, and heterocyclic so it has potential to be processed into synthetic fuel. Coal tar is mostly composed of C, H, and O, and also lower concentration of S and N [1]. The aromatics and heteroatoms (S and N) contained in coal tar must be changed and the molecular weight and viscosity should be lowered. To lower the molecular weight, viscosity, and aromatics content, coal tar must be processed by hydrogenation and cracking. Hydrogenation and cracking process requires a catalyst that has dual functions, metal component as the hydrogenation catalyst and acid component as the cracking catalyst [2]. One of the hydrogenation metals most commonly used in the hydrocracking process are Co and Mo. While the most common catalyst used in catalytic cracking and hydrocracking is Y zeolite. Co-Mo/Y zeolite is expected to have high activity in coal tar hydrocracking process [3].

Wang, et al [4] had did a research using zeolite catalyst, that reaction initially took place at high temperature, but by using catalyst the reaction could take place at lower temperature. Research by Lestari et al [5] about the effect of Ni and Mo metal showed that catalyst with Ni and Mo required higher specific surface area to produce greater distribution of Mo. Ramadan and Friandani [6] studied the effect of Co and Mo metal addition on Y zeolite where the higher concentration of the metal, the higher values of catalyst acidity was if supported by equitable distribution. Therefore this research impregnated Co and Mo into Y zeolite to convert coal tar into liquid fuels and not Ni metal which is more expensive and requires greater specific surface area.

The effect of impregnation temperature and time need to be considered to determine the best preparation of Co-Mo/Y zeolite condition which can be used as a hydrocracking

catalyst of coal tar. This has encouraged research to determine the effect of impregnation condition Co and Mo into Y zeolite in coal tar hydrocracking into liquid fuel.

The purpose of this study was to determine the catalyst preparation Co-Mo / zeolite Y with impregnation method, knowing the characteristics of the catalyst Co-Mo / zeolite Y, determine the effect of temperature and time of impregnation on the preparation of the catalyst Co-Mo / zeolite Y for the conversion process coal tar, and determine the best conditions in the manufacture of Co-Mo / zeolite Y for the conversion process coal tar.

EXPERIMENTAL SECTION

Materials

The materials used are coal tar from PT. Sango Ceramic Indonesia, Y zeolite and ZSM-5 from PT. Zeolyst International, $\text{Co}(\text{NO}_3)_2 \cdot 6\text{H}_2\text{O}$ from Merck, $(\text{NH}_4)_6\text{Mo}_7\text{O}_{24} \cdot 4\text{H}_2\text{O}$ from Merck, ammonia from Merck, hydrogen gas from PT. Aneka Gas, and distilled water from Chemical Process Laboratory Department of Chemical Engineering Universitas Diponegoro.

Instrumentation

The functional group of the catalysts was analyzed by Fourier Transform-Infra Red (FTIR) in Technology Laboratory Separation Department of Chemical Engineering, University of Diponegoro. Stability catalyst structure by X-Ray Diffraction (XRD) Laboratory Integrated Diponegoro University, and the total acidity method adsorbs ammonia gravimetrically at Laboratory of Basic Chemical Engineering Department Chemistry University of Diponegoro. Liquid fuel product analyzed using Gas Chromatography at Organic and Bioorganic Laboratory Department of Chemistry Universitas GadjahMada in standard gasoline.

Procedure

Preparation and Characterization of Catalyst

Preparation of Co-Mo/Y zeolite catalyst used impregnation method. Five grams of Y zeolite, cobalt metal obtained from $\text{Co}(\text{NO}_3)_2 \cdot 6\text{H}_2\text{O}$ 0.985 grams, and molybdenum metal from $(\text{NH}_4)_6\text{Mo}_7\text{O}_{24} \cdot 4\text{H}_2\text{O}$ 4.34 grams, dissolved in 50 ml of distilled water. The solution was then heated in accordance temperature variables during the time variables. The samples then filtered using a vacuum pump until no more water was dripping. Further drying was done in the oven with temperature of 100 °C for 24 hours. Dried samples then calcined at

temperature of 550°C for 3 hours. The functional group of the catalysts was analyzed by FTIR. Stability catalyst structure was analyzed by X-Ray Diffraction. The total acidity method adsorbs ammonia gravimetrically method.

Catalys Testing and Analysis of Product

Hydrocracking process used a 1000 ml three-neck flask size as a place of coal tar evaporation at 300°C, with hot plate and magnetic stirrer. Hydrogen gas was used as the driving medium and also to assist the cracking process. The reactor used was a pipe reactor in which ceramic and glasswool used to support the catalyst to remain in its position. There were temperature indicator and heater in the reactor so that the temperature reaction was maintained at 350°C. Seven grams catalyst, consisted of 5 grams catalyst which had been prepared (Co-Mo/Y zeolite) and 2 grams ZSM-5 catalyst, put in the pipe reactor. Cooling used a screw cooler and ethylene glycol as coolant. Hydrocracking process was carried out for 1 hour since the first droplet. Liquid fuel product then analyzed using Gas Chromatography at Organic and Bioorganic Laboratory Department of Chemistry Universitas Gadjah Mada in standard gasoline.

Experimental Design

Method used for achieving the goals is response surface methodology central composite rotary design. In this study, there are two independent variables, impregnation temperature and time. Limitation of independent variables is shown in **Table 1** processed using STATISTICA software.

RESULTS AND DISCUSSION

Characterization Using Analysis of FTIR (Fourier Transform-Infra Red)

Catalyst characterization is intended to determine whether Co and Mo had impregnated into Y zeolite. **Figure 1** is an IR spectrum which shows the characteristic of Y zeolite and Co-Mo/Y zeolite. From the interpretation of the spectrum **Figure 1** indicates that impregnation Co and Mo at impregnation temperature variables and time and didn't cause changes in the structure of catalyst. Preparation of catalyst with impregnation method aims to elicits metal which can expand the surface active site catalyst.

When compared with standard spectra of Y zeolite, the characterization results of Co-Mo/Y zeolite catalyst showed that the ten samples had frequency or wave number shifted at 1199.87 cm^{-1} and 1067.88 cm^{-1} , which indicates the range of O-Si-O and O-Al-O [7].

Presumably this shift is caused by a reduction of Al atoms in the framework become Al non-framework due to dealuminated at catalyst calcination [7]. At 200°C, protons in zeolite have high mobility and at 550° C are separated as water to form the Lewis site, can be described in **Figure 2**.

The presence of water vapor will constantly strengthen the structure stability of Lewis site structure which is not stable, the result is called "actual Lewis sites" which can be seen in **Figure 3**.

Peak in the standard spectrum of Y zeolite at 1400.62 cm^{-1} is present while not in ten samples. According to Coates [7], the peak indicates -OH. The cause of the loss of is by hydrothermal process when the Co-Mo/Y zeolite catalyst calcination. There is also a peak at 829.49 cm^{-1} on spectrum of Y zeolite standard which indicates C-H [7]. Meanwhile the spectrums of Co-Mo/Y zeolite have a shift at that wave numbers thought caused by impregnation of Co and Mo. Also, there is appearance of two peaks near wave numbers of 902 cm^{-1} and 945 cm^{-1} while there is none in spectra standard of Y zeolite before. The area indicates asymmetrical strain O-Si-O and O-Al-O [7]. The appearance strengthens the presumption of Co and Mo had impregnated into catalyst accordance impregnation temperature and time variables.

Effect of Impregnation Temperature and Time To Co-Mo/Y Zeolite Crystallinity

It can be observed in **Figure 4** that the intensity of the ten of Co-Mo/Y zeolite catalysts' diffractogram is decreasing. Catalyst crystallinity is calculated by comparing the intensity of 8 strongest peaks Co-Mo/Y zeolite with Y zeolite. So we get the data as shown in **Table 2**.

Catalyst crystallinity decreased when Co and Mo were impregnated. This is probably caused by the impregnation of Co and Mo into Y zeolite. However, the ten catalysts have similar pattern with Y zeolite, so it can be said that the crystallinity of Y zeolite was not damaged after impregnating Co and Mo [8]. This is consistent with the results of FTIR analysis where the graph shows that the structure does not change after impregnating Co and Mo.

Based on **figure 5 (a)**, at the same impregnation time can be seen that the higher the temperature, the higher degree of catalyst crystallinity is. According to Krichko [9], the increasing of impregnation temperature may increase the mobility of the chains bond in the catalyst, which may occur in the area of the crystal structure. High mobility led to the composition of the chain becomes thight and thus expand the areas of crystal structure. However, there are deviations in the chart, which is a decrease of crystallinity as the temperature rises at impregnation time of 20 minutes and temperature of 60°C.

Figure 5 (b) indicates that at the same impregnation temperature, there is an increase in crystallinity as the longer impregnation time. The decline occurred because the metal deposits on the catalyst samples that cover the surface of the pores so can change the characteristics of the zeolite crystal and causes a decrease in the intensity of the curve [10]. The longer impregnation time causes Co and Mo are more impregnated, and the catalyst crystallinity has decreased.

Effect of Impregnation Temperature and Time To Total Acidity

According to **figure 6** can be seen that increasing of total acidity occurs after impregnating Co and Mo into Y zeolite. The measured acidity of catalyst is the sum total of Bronsted acid and Lewis acid. In the catalysts impregnated metal, metals donate the amount of the acidic Lewis. This is consistent with the analysis results showed an upward trend in the acidity of the catalyst with the addition of Co and Mo into Y zeolite [10,11].

At the same impregnation time, it can be seen in **Figure 7 (a)** that the increasing of impregnation temperature increases the total acidity at 10 minutes and decreases at 30 minutes. While at 20 minutes decreasing occurs then increasing. To find out the cause of this phenomenon is expected for further Co-Mo/Y zeolite catalyst characterization.

Based on **figure 7 (b)**, at same impregnation temperature can be seen that the longer impregnation time, the total acidity of the catalyst is increased. This is likely due to the longer time, the more ammonia is absorbed by Co-Mo/Y zeolite. However, a deviation occurs at 45°C and 20 minutes. The presumption of this deviation is caused by uncompletely distribution of ammonia adsorption.

Effect Impregnation Temperature and Time To Liquid Fuel Product

According to **figure 8 (a)**, at the same impregnation temperature, increasing in time causes increasing in yield. One of the factors that may affect the amount of the product is the catalyst, so the catalyst characteristic, ie acidity and crystallinity, effect on the product formed. This is consistent with the initial discussion in which the increase in time lead to increases in acidity. High acidity catalyst causes catalyst performance increases. Therefore, the reaction formation of liquid fuels from coal tar faster and produce more yield.

In **Figure 8 (b)**, increasing in yield occurs as the temperature rises at the same time. This is influenced by increasing in crystallinity as temperature rises. The high percentage of crystallinity indicates that the catalyst structure did not change significantly and tend to be more stable, so the ability of the catalyst can increase and accelerate the formation of

reaction product. Data crystallinity and the concentration of the gas in the GC product analysis results are shown in **Table 3**.

From **table 3**, it is obtained relationship between Co-Mo/Y zeolite crystallinity at impregnation temperature and time variables with the concentration of the gas contained in coal tar products of hydrocracking.

According to **figure 9** it can be seen that the higher percentage of Co-Mo/Y zeolite crystallinity, the greater the concentration of the fuel produced. Crystallinity is a very important factor influencing the nature of the catalyst. High crystallinity indicate that catalyst is free from impurities and physical properties (catalytic properties of high, stable at high temperatures and extensive porosity) are not disturbed [10,12]. As a result, performance can be run optimally, so that produce more product.

Analysis of Liquid Fuel

GCMS analysis obtained peak gasoline in the range of 1.3-19.9 minutes and GC analysis obtained 10 products with different peak. Concentration gasoline on 10 products can be determined by calculating the concentration peak GC results were in the range of gasoline when GCMS results. Commercial gasoline sample is obtained from gas station where GCMS analysis indicates that the components contained in the gasoline have a number of chains of carbon atoms ranging from C_2 - C_{10} .

Consideration of the suitable using of hydrocarbons for gasoline is based on the volatility and octane number. In general, gasoline has a component of C_5 - C_8 , but some are up to C_9 or more [13]. In the range of C_5 - C_8 , it contained aromatic isomers with branched-chain or cyclic good in the use of gasoline as a motor fuel. This is consistent with the results of GC analysis of 10 products in Table 4 that most of the products contain C_8 , C_9 and C_{10} . **Table 4** shows that the best impregnation condition to get the most yield, 1.35 ml of gasoline, is by impregnation at temperature of 55°C and 30 minutes.

CONCLUSION

Catalyst characterization results indicate that Co and Mo had impregnated into Y zeolite. Impregnation of Co and Mo at impregnation temperature and time variables did not damage the structure of Y zeolite catalyst and increased the value of Y zeolite total acidity. Increasing in impregnation temperature caused catalyst crystallinity and acidity as well as the yield of gasoline rised. Meanwhile, the longer the impregnation time, the more reduced the catalyst crystallinity but the catalyst total acidity and yield increased. GC analysis

showed that the products in the time range of gasoline contained components of C₈, C₉, and C₁₀. To get the best crystallinity Co-Mo/Y zeolite catalyst was by impregnating at temperature range of 40-50°C and time of 5-20 minutes. Meanwhile, to get the best yield, ie 1.35 ml of gasoline, was by using a catalyst with impregnation method at temperature of 55°C and 30 minutes

ACKNOWLEDGEMENTS

The authors gratefully acknowledge the financial support from Diponegoro University DIPA Number 214-05/UN7.5.1/PG/2015 and support from PT. Sango Ceramic Indonesia for Feedstock Coal Tar.

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Table 1 Matrix of experimental design

Run	Temperature (°C)	Time (minutes)
1	35	10
2	35	30
3	55	10
4	55	30
5	31	20
6	59	20
7	45	6
8	45	34
9	45	20
10	45	20

Table 2 Co-Mo/Y zeolite crystallinity

Run	Crystallinity (%)
1	23,84
2	17,39
3	36,49
4	17,73
5	25,35
6	23,31
7	49,84
8	30,86
9	46,53
10	43,92

Table 3. Crystallinity and concentration of gasoline produced by Co-MO/Y zeolite

Run	Crystallinity (%)	Concentration of Gasoline (%)
1	23,84	36,54
2	17,39	37,23
3	36,49	71,19
4	17,73	89,80
5	25,35	44,65
6	23,31	89,12
7	49,84	89,30
8	30,86	56,81
9	46,53	91,51
10	43,92	83,93

Table 4. Yield and composition of C₈, C₉, and C₁₀ in gasoline

Run	Yield of Gasoline (ml)	Composition (%)		
		C ₈	C ₉	C ₁₀
1	0,40	0,00	90,31	9,69
2	0,41	0,00	0,00	100,00
3	0,71	9,13	9,11	81,77
4	1,35	2,78	1,53	95,69
5	0,54	0,00	56,30	43,70
6	1,16	0,00	95,86	4,14
7	0,80	11,60	7,73	80,67
8	1,02	0,00	0,00	100,00
9	1,10	2,18	57,96	39,86
10	0,76	2,92	0,00	97,08

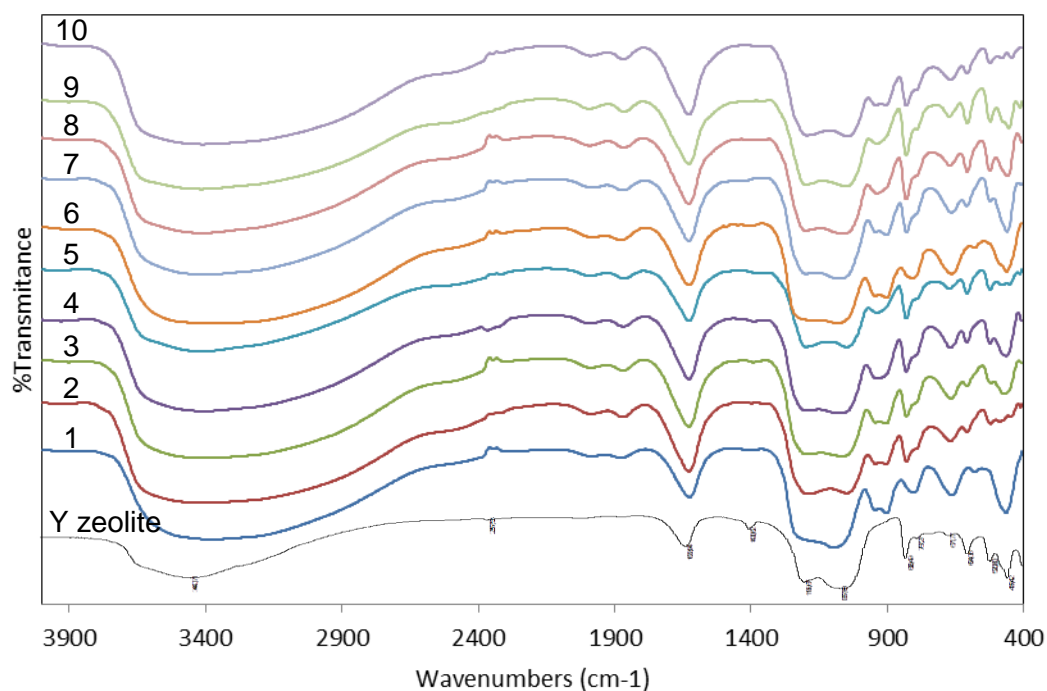


Figure 1 FTIR spectra of Y zeolite Y and Co-Mo/Y zeolite

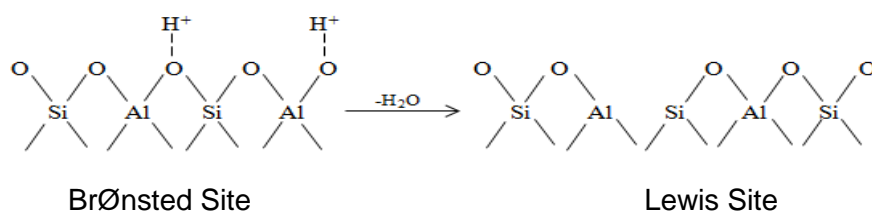


Figure 2 The process of release of a proton from Brønsted site into Lewis site

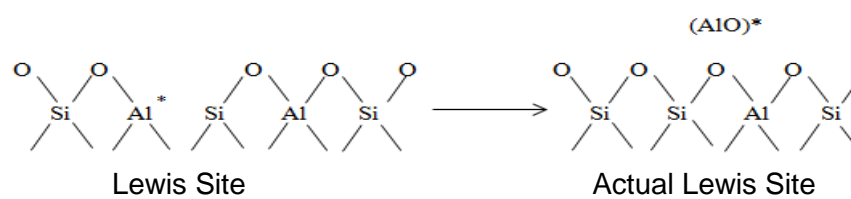


Figure 3 The process of structural stability from Lewis site into actual Lewis site

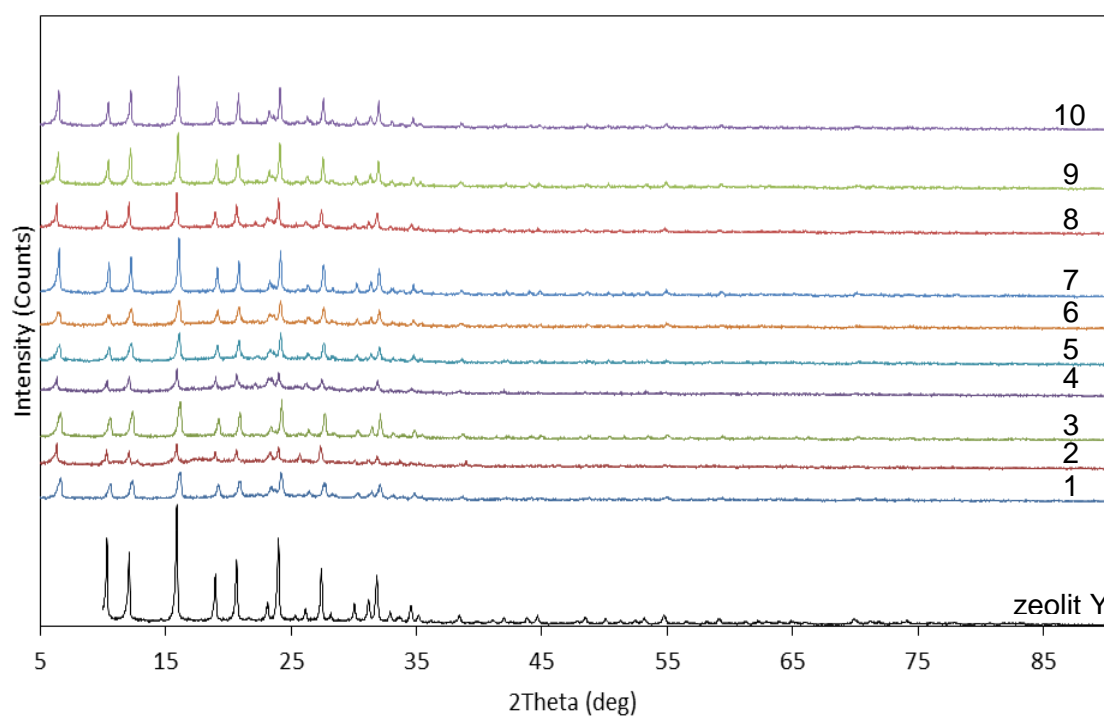


Figure 4 XRD diffractogram of Y zeolite and Co-Mo/Y zeolite

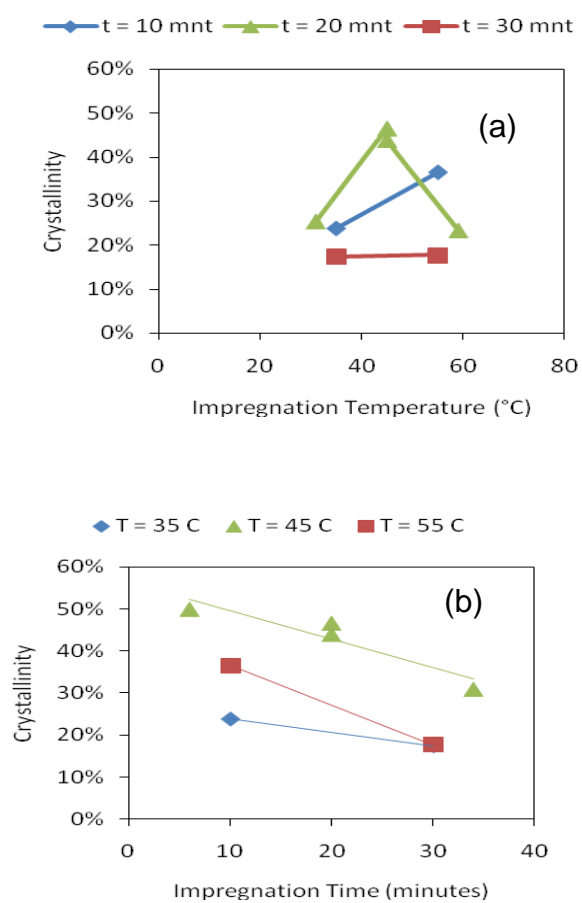


Figure 5. Relationship of impregnation (a) temperature and (b) time with crystallinity

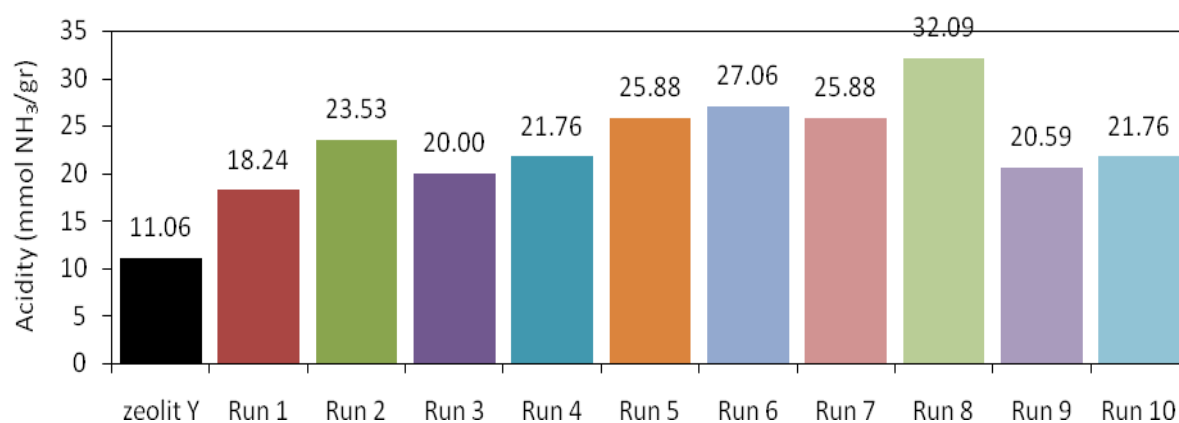


Figure 6 Value of Y zeolite and Co-Mo/Y zeolite total acidity

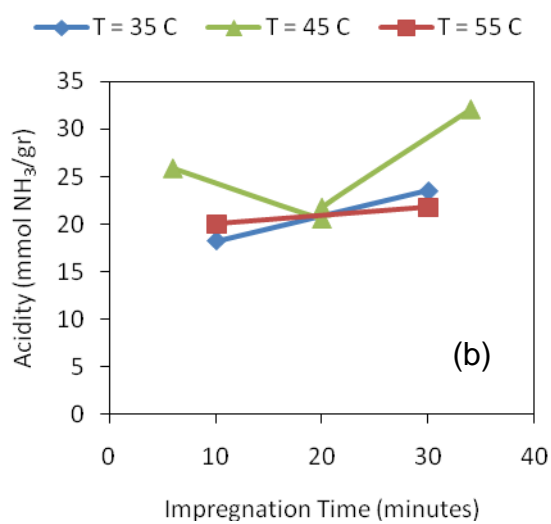
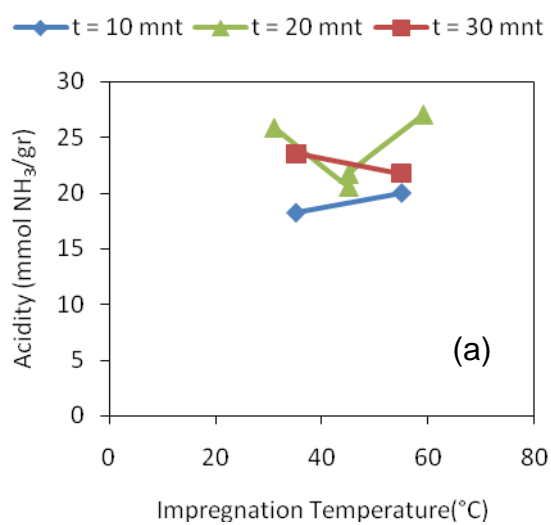


Figure 7 Relationship of impregnation (a) temperature and (b) time with acidity

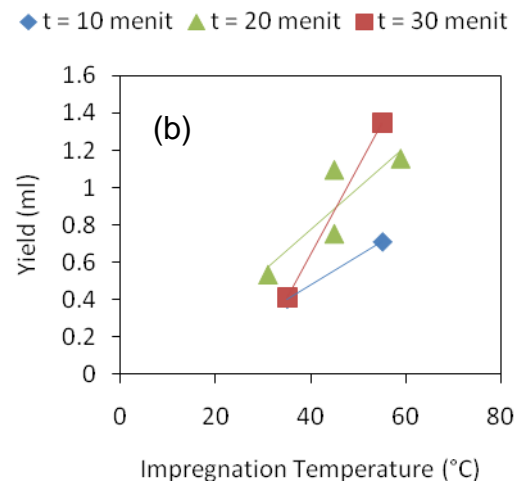
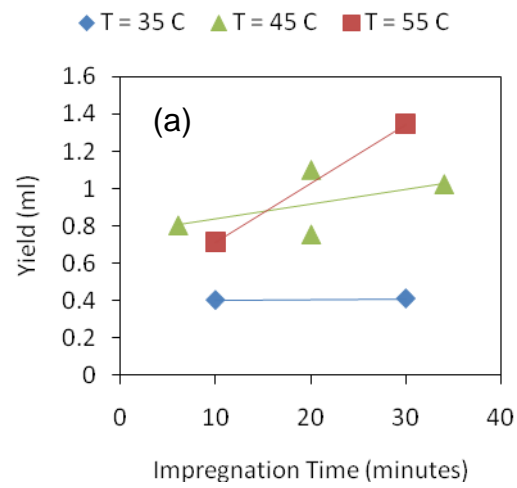


Figure 8 Relationship of impregnation (a) time and (b) temperature with yield

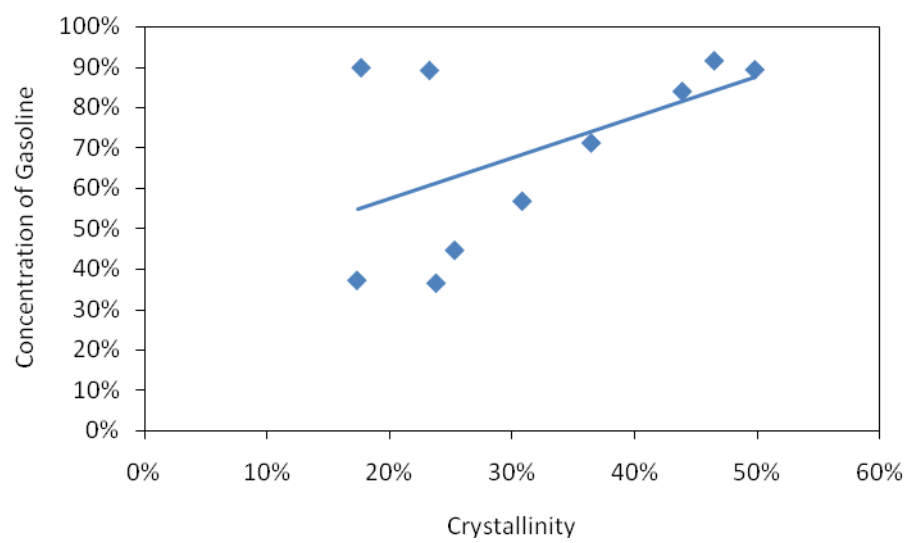


Figure 9 Relationship between crystallinity and concentration of gasoline

REVISION NOTE

REVIEWER A		
No	Comments	Answer
1	The unit in tables and figures are not consistently written The English used in this paper is rather below standard, so huge efforts are should be taken to improve it.	Already improved
2	It seems that the paper was prepared under google translator rather than being checked by a native speaker.	Already English improved
3	In addition, many typographical errors are found throughout the paper. Some revisions are necessary to improve the readability of this paper. Author should obey the comments below and responding them accordingly.	Already corrected of typographical and improved of readability.
4	Affiliation: University of Diponegoro should be Diponegoro University	Already corrected
Abstract:Page 1.		
1	The aims of the research are unclear.	Already corrected to clearly
2	The statement is confusing Is it true that the coal tar is reacted with the catalyst?	Already corrected the statement that the coal tar is reacted with the catalyst.
3	Did you mean the “hydrogen introduction? Rather than hydrogen flowing?	Already corrected “Hydrogen gas (as a reactant)”
4	This phrase “Characteristic analysis showed that Co and Mo had impregnated into Y zeolite, as well as it made no change of catalyst’s structure and increased the total acidity.	<i>Characteristic analysis showed that Co and Mo had impregnated into Y zeolite, as well as it made no change of catalyst’s structure and increased the total acidity.</i>
5	The higher the impregnation temperature was, the higher catalyst crystallinity, total acidity, and yield were.	<i>The higher of impregnation temperature was increased the catalyst crystallinity, total acidity, and yield of gasoline.</i>
6	The longer impregnation took time, the more crystallinity reduced but total acidity and yield increased.	<i>The longer impregnation time</i>

		<i>was reduced crystal linity value but total acidity and yield were increased.</i>
7	GC analysis showed that product in the time range of gasoline contained C8, C9, and C10.” is even more confusing to readers. Please rephrase and make it more readable.	<i>GC analysis showed that products included into the gasoline product (C₈, C₉, and C₁₀).</i>
Introduction:		
1	Please give reference for the first paragraph of introduction	Already give the reference
2	Please rephrase the last two paragraphs of this section. They are unclear and could develop confusion over the readers	Already rephrase and corrected to clearly
Material and Method:		
1	What do the authors mean by “Technology Laboratory Separation Department of Chemical Engineering”	Revision: Chemical Process Laboratory Department of Chemical Engineering, Diponegoro University.
2	What do the authors mean by “the total acidity method adsorbs ammonia gravimetrically at Laboratory of Basic Chemical Engineering Department Chemistry University of Diponegoro.”.	Revision: method adsorbs ammonia gravimetrically was done at Laboratory of Basic Chemical Engineering Department Chemistry University of Diponegoro
3	Please rephrase the “procedure for catalyst preparation”	Already rephrase
Catalys Testing and Analysis of Product (please make correction)		
1	Are the authors sure that the hot plate could bring the temperature of the three necked flash to about 300oC? It is rather ridiculous.	Hydrocracking process used a 1000 ml three-neck flask size as a place of coal tar (100 ml), for evaporated using

		a hot plate (300°C), above a hot plate there is oil as media warm up and magnetic stirrer.
2	Please also rephrase the section "Catalys Testing and Analysis of Product"	Already rephrase.
Results and Discussion:		
1	Please give detail discussion of your results rather than just reporting the value of data.	Already explanation of data.
2	It is better to express yield in (amount of product/amount of feed or multiplied it by 100 to get % yield). Reporting yield in the form of mL is not appropriate because we do not know the change of the reactant into the product. See. Figure 8	Already correction.
3	The experimental data in Figure 8 and Figure 9 are not linear, but the authors forced them to be linear. Why?	Figure 8 (now as Figure 9) is changed to be linear. Figure 9 (now as Figure 10) is linear, without crystallinity at 17.73 and 23.31%.
4	Some figure still comes with Bahasa Indonesia (menit)	Already correction.
Conclusion:		
1	Please make your conclusion in brief, clear and concise.	Already correction and improve.
2	The current conclusion is unclear and confusing.	Already correction and improve.
References:		
1	Please write the literature cited following the guideline that may be obtained from the BCREC website	Already correction and improve.
REVIEWER B		
Page	Comments	Answer
1	[1] Please check English Grammar throughout the texts (all sentences) in this paper, sentence by sentence, carefully. I suggest you to use Grammarly software	Already correction and improve.
1	[2] Keyword must be specific and represent this research. Do not use too general words	Already correction and improve.
1	[3] Please delete this section	Already delete.
1	[4] Please delete this section	Already delete.
2	[5] Grammar????? "...had did.."	Already correction.
3	[6] In the end of Introduction section, authors MUST put gap analysis statements or novelties of this paper, what is	Already correction and improve.

	unique finding of this paper compare to previous other research. Thus put statement of objectives of this paper	
3	[7] Please state the purity of chemicals	Already state the purity of chemicals.
3	[8] Brand and model of equipment???	Already explain of brand and model.
4	[9] Grammatical error????	Already correction.
4	[10] Please provide experimental rig schematically	Already provide experimental rig figure (1)
4	[11] Please state range of variables in this experiment. Please state table of experimental design.	Already state of variables.
6	[12] So many typo and spelling errors. Please check them throughout text carefully	Already correction.
7	[13] Avoid a paragraph containing only a sentence. A paragraph should contains of an idea dan its explanation	Already more explanation.
7	[14] So many grammatical errors	Already correction.
7	[15] What is relation between liquid fuels yield or performance and the catalyst characterization???	Already more explanation.
8	[16] So many grammatical error	Already correction and improve.
8	[17] Please type references as Author Guidelines (URL: http://ejournal2.undip.ac.id/index.php/bcrec/pages/view/authorguide)	Already correction and follow guidelines
9	[18] Comma or dot???	Already change comma to dot.
9	[19] Comma or dot???	Already change comma to dot.
10	[20] Comma or dot???	Already change comma to dot.
11	[21] Please only depict on 2 theta ranges of 5 to 45 only	Already depict on 2 theta ranges 5-45 only

2nd Revision Note File

No	Comment	Correction
1	DO NOT shorten the last names !!	Already correction.... GIVENI CHRISTINA SILAEN, RESTI NUR UTAMI
2	Please improve English grammar in this sentences	The cooling system used a screw condenser and ethylene glycol (as coolant). Time of reaction is 1 hour. Then, analyze of liquid fuel product carried by....
3	In this method, author told that the design experiment of RSM using STATISTICA has been used, BUT in the results there is no optimization results using the RSM, no contour, no surface simulation???	Already delete
4	Please write the formula and method to calculate this cristalinity in %.	Already write in page number 6...The crystallinity of the zeolite can be calculated from X-ray diffractograms and is expressed according to [20].
5	% here is concentration or percentage of composition???? As far as I know that concentration is in mol/volume??	Already change ... concentration to composition
6	Comma or dot?	Already change ... comma to dot
7	Yield, usually presented in “%” based the reactant and product. Not in ml.	Already change...ml to %
8	X-axis and unit ????	Figure 6(a) already correction of X-axis and unit.
9	X-axis and unit ????	Figure 6(b) already correction of X axis and unit.
10	X-axis and unit ????	Figure 8(a) already correction of X-axis and unit.
11	X-axis and unit ????	Figure 8(b) already correction of X-axis and unit.
12	X-axis and unit ????	Figure9(a) already correction of X-axis and unit.
13	X-axis and unit ????	Figure9(b) already correction of X-axis and unit.
14	Unit of x-axis and y-axis labels?	Figure 10 already correction of Unit of x-axis and y-axis labels?